

**REMARKS**

Applicants have amended the specification to cross reference the parent applications of which this application is a continuation of a pending U.S. application, which is a continuation of a PCT application designating the United States which itself is a continuation of a U.S. provisional application.

Applicants have also amended claims 3, 5-6, 8, 10-11 and 13-15 and canceled claims 17-30 in order to reduce the filing fee by deleting the multiple dependencies and additional independent claims. Applicants retain the right to reintroduce any subject matter canceled by the present Amendment at any time during the prosecution of this application or any continuation or divisional thereof in the United States.

The present application is a continuation application and the prior art cited in the parent applications should be taken into consideration in the present application. In accordance with MPEP §2001.06(b) no copies of the prior art in the parent applications are submitted herewith. The reference cited forms from the parent applications are submitted herewith for the convenience of the Examiner. In accordance with MPEP §609, a Form 1449 listing these references is also submitted herewith. Confirmation that the prior art cited in the parent applications has been considered in the next Official Action is most respectfully requested.

In view of the above amendments to the claims an early and favorable action on the merits is now in order and is most respectfully requested.

Respectfully submitted,  
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**Marked-Up Version Showing Changes Made**

**IN THE CLAIMS:**

Please replace claims 3, 5-6, 8, 10-11 and 13-15 with the following amended claims.

3(Amended). A method as claimed in claim 1 [or claim 2] wherein said imaging procedure is a gradient echo or echo planar imaging procedure.

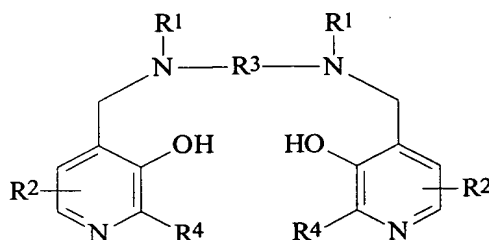
5(Amended). A method as claimed in claim 3 [or claim 4] wherein said imaging procedure is one in which TI (inversion time) is 100 to 800 msec, TR (repetition time) is 2000 msec and TE (echo time) is less than 20 msec.

6(Amended). A method as claimed in [any preceding claim] claim 1 wherein said manganese complex or salt thereof is administered at a dosage of 0.005 to 0.2 mmol/kg bodyweight.

8(Amended). A method as claimed in [any preceding claim] claim 1 wherein said manganese complex is a manganese chelate complex having a  $K_a$  value of from  $10^7$  to  $10^{25}$ .

10(Amended). A method as claimed in claim 8 [or claim 9] wherein said chelate has a  $K_a$  value smaller by a factor of at least  $10^3$  than the  $K_a$  value of the corresponding ferric ( $Fe^{3+}$ ) chelate.

11(Amended). A method as claimed in [any one of claims 8 to 10] claim 8 wherein said manganese chelate comprises a chelating compound of formula I:



(I)

or a salt thereof

(wherein in formula I

each  $R^1$  independently represents hydrogen or  $-\text{CH}_2\text{COR}^5$ ;

$R^5$  represents hydroxy, optionally hydroxylated alkoxy, amino or alkylamido;

each  $R^2$  independently represents a group  $\text{XYR}^6$ ;

X represents a bond, or a  $\text{C}_{1-3}$  alkylene or oxoalkylene group optionally substituted by a group  $R^7$ ;

Y represents a bond, an oxygen atom or a group  $\text{NR}^6$ ;

$R^6$  is a hydrogen atom, a group  $\text{COOR}^8$ , an alkyl, alkenyl, cycloalkyl, aryl or aralkyl group optionally substituted by one or more groups selected from  $\text{COOR}^8$ ,  $\text{CONR}^8$ ,  $\text{NR}^8$ ,  $\text{OR}^8$ ,  $=\text{NR}^8$ ,  $=\text{O}$ ,  $\text{OP}(\text{O})(\text{OR}^8)\text{R}^7$  and  $\text{OSO}_3\text{M}$ ;

$R^7$  is hydroxy, an optionally hydroxylated, optionally alkoxyated alkyl or aminoalkyl group;

$R^8$  is a hydrogen atom or an optionally hydroxylated, optionally alkoxyated alkyl group;

M is a hydrogen atom or one equivalent of a physiologically tolerable cation;

$R^3$  represents a  $\text{C}_{1-8}$  alkylene group, a 1,2-cycloalkylene group, or a 1,2-arylene group; and

each  $R^4$  independently represents hydrogen or  $\text{C}_{1-3}$  alkyl).

13(Amended). A method as claimed in claim 11 [or claim 12] wherein in formula I,  $R^3$  is ethylene and each group  $R^1$  represents  $-\text{CH}_2\text{COR}^5$  in which  $R^5$  is hydroxy.

15(Amended). A method as claimed in [any one of claims 8 to 10] claim 8 wherein said chelate complex is a complex of a linear, branched or macrocyclic chelant selected from polyaminopolycarboxylic acid chelants and carboxylic acid derivatives thereof.